Amendments to the claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

1. (Currently amended) A compound of formula (I):

(I)

wherein

A is a fused 5-membered heteroaryl ring containing up to two heteroatom independently selected from oxygen, nitrogen or sulfur, optionally substituted by up to two substituents independently selected from C_{1-6} alkyl, $-(CH_2)_k-C_{3-7}$ cycloalkyl, halogen, -CN, trifluoromethyl, $-(CH_2)_kOR^3$, $-(CH_2)_kCO_2R^3$, $-(CH_2)_kNR^3R^4$, $-(CH_2)_kCONR^3R^4$, $-(CH_2)_kNHSO_2R^3$, $-(CH_2)_kSO_2(CH_2)_mR^5$, a 5-or 6-membered heterocyclyl ring containing nitrogen optionally substituted by C_{1-2} alkyl or $-(CH_2)_kCO_2R^3$, and a 5-membered heteroaryl ring optionally substituted by C_{1-2} alkyl; or

A is a fused 5-membered heteroaryl ring containing up to two heteroatom independently selected from oxygen, nitrogen or sulfur substituted by $-B^1R^6$, and A is optionally further substituted by one substituent selected from $-OR^7$, halogen, trifluoromethyl, -CN, $-CO_2R^7$ and C_{1-6} alkyl optionally substituted by hydroxy; or

A is a fused 5-membered heteroaryl ring containing up to two heteroatom independently selected from oxygen, nitrogen or sulfur substituted by -(CH₂)_nheterocyclyl wherein the heterocyclyl is a 5- or 6-membered heterocyclic ring containing one or two heteroatoms independently selected from oxygen, sulfur and nitrogen optionally substituted by up to two substituents independently selected from oxo, C_{1-6} alkyl, OR^7 , - NR^7R^8 and - $CONR^7R^8$, and A is optionally further substituted by one substituent selected from - OR^7 , halogen, trifluoromethyl, -CN, - CO_2R^7 and C_{1-6} alkyl optionally substituted by hydroxy; or

A is a fused 5-membered heteroaryl ring containing up to two heteroatom independently selected from oxygen, nitrogen or sulfur substituted by -(CH₂)_qaryl or -(CH₂)_qheteroaryl wherein the aryl or heteroaryl is optionally substituted by one or more substituents independently selected from oxo, C₁₋₆alkyl, halogen, -CN, trifluoromethyl, -OR⁹, -(CH₂)_rCO₂R¹⁰, -NR⁹R¹⁰, -(CH₂)_rCONR⁹R¹⁰, -NHCOR⁹, -SO₂NR⁹R¹⁰, -NHSO₂R⁹ and -S(O)₈R⁹, and A is optionally further substituted by one substituent selected from -OR⁷, halogen, trifluoromethyl, -CN, -CO₂R⁷ and C₁₋₆alkyl optionally substituted by hydroxy;

R¹ is selected from methyl and chloro;

R² is selected from -NH-CO-R¹¹ and -CO-NH-(CH₂)_t-R¹²;

 R^3 is selected from hydrogen, $C_{1\text{-}6}$ alkyl optionally substituted by up to two OH groups, -(CH₂)_k-C₃₋₇cycloalkyl, -(CH₂)_kphenyl optionally substituted by R^{13} and/or R^{14} and -(CH₂)_kheteroaryl optionally substituted by R^{13} and/or R^{14} ,

R⁴ is selected from hydrogen and C₁₋₆alkyl, or

R³ and R⁴, together with the nitrogen atom to which they are bound, form a 5-or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R¹⁵;

 R^5 is selected from $C_{1\text{-}6}$ alkyl optionally substituted by up to three halogen atoms, $C_{2\text{-}6}$ alkenyl optionally substituted by phenyl, $C_{3\text{-}7}$ cycloalkyl, heteroaryl optionally substituted by up to three R^{13} and/or R^{14} groups, and phenyl optionally substituted by R^{13} and/or R^{14} :

 R^6 is a C_{3-6} alkyl group substituted by at least two substituents independently selected from -OR¹⁶, -NR¹⁶R¹⁷, -CO₂R¹⁶, -CONR¹⁶R¹⁷, -NHCOR¹⁶ and -NHSO₂R¹⁶;

 ${\sf R}^7$ and ${\sf R}^8$ are each independently selected from hydrogen and ${\sf C}_{1\text{-}6} {\sf alkyl};$

 $\rm R^9$ is selected from hydrogen, -(CH₂)_u-C₃₋₇cycloalkyl, -(CH₂)_uheterocyclyl, -(CH₂)_uaryl, and C₁₋₆alkyl optionally substituted by up to two substituents independently selected from -OR¹⁸ and -NR¹⁸R¹⁹,

 R^{10} is selected from hydrogen and $C_{1\text{--}6}$ alkyl, or

 R^9 and R^{10} , together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R¹⁵;

 R^{11} is selected from hydrogen, $C_{1\text{-}6}$ alkyl, -(CH₂)_t- $C_{3\text{-}7}$ cycloalkyl, trifluoromethyl, -(CH₂)_vheteroaryl optionally substituted by R^{20} and/or R^{21} , and -(CH₂)_vphenyl optionally substituted by R^{20} and/or R^{21} ;

 $\rm R^{12}$ is selected from hydrogen, C $_{1-6}$ alkyl, C $_{3-7}$ cycloalkyl, -CONHR 22 , phenyl optionally substituted by R 20 and/or R 21 , and heteroaryl optionally substituted by R 20 and/or R 21 ;

 R^{13} and R^{14} are each independently selected from halogen, -CN, trifluoromethyl, nitro, C_{1-6} alkyl, C_{1-6} alkoxy, -CONR 22 R 23 , -COR 24 , -CO2R 24 , and heteroaryl, or

 R^{13} and R^{14} are linked to form a fused 5-membered heterocyclyl ring containing one heteroatom selected from oxygen, sulfur and N-R¹⁵, or a fused heteroaryl ring;

R¹⁵ is selected from hydrogen and methyl;

 $R^{16},\,R^{17},\,R^{18}$ and R^{19} are each independently selected from hydrogen and $C_{1\text{-}6}alkyl;$

 R^{20} is selected from C_{1-6} alkyl, C_{1-6} alkoxy, - $(CH_2)_t$ - C_{3-7} cycloalkyl, - $CONR^{22}R^{23}$, -NHCOR²³, halogen, -CN, - $(CH_2)_w$ NR²⁵R²⁶, trifluoromethyl, phenyl optionally substituted by one or more R²¹ groups, and heteroaryl optionally substituted by one or more R²¹ groups;

 R^{21} is selected from C $_{1\text{-}6}$ alkyl, C $_{1\text{-}6}$ alkoxy, halogen, trifluoromethyl, and -(CH $_2)_wNR^{25}R^{26};$

 R^{22} and R^{23} are each independently selected from hydrogen and $C_{1\text{-}6}$ alkyl, or R^{22} and R^{23} , together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R¹⁵, wherein the ring may be substituted by up to two $C_{1\text{-}6}$ alkyl groups;

 R^{24} is C_{1-6} alkyl;

 R^{25} is selected from hydrogen, $C_{1\text{-}6}$ alkyl and - $(CH_2)_t$ - $C_{3\text{-}7}$ cycloalkyl optionally substituted by $C_{1\text{-}6}$ alkyl,

 R^{26} is selected from hydrogen and C_{1-6} alkyl, or

 R^{25} and R^{26} , together with the nitrogen atom to which they are bound, form a 5- or 6-membered heterocyclic ring optionally containing one additional heteroatom selected from oxygen, sulfur and N-R¹⁵;

 R^{27} is hydrogen or $C_{1\text{-}6}$ alkyl;

 B^{1} is selected from a bond, oxygen, NH and $S(O)_{x}$;

X and Y are each independently selected from hydrogen, methyl and halogen;

Z is selected from halogen, C₁₋₆alkyl and -OR²⁷;

k, m and w are each independently selected from 0, 1, 2 and 3;

n, q, r, s, t and x are each independently selected from 0, 1 and 2; and u and v are each independently selected from 0 and 1;

or a pharmaceutically acceptable salt thereof.

- 2. (original) A compound according to claim 1 wherein A is a fused 5-membered heteroaryl ring containing up to two heteroatoms independently selected from oxygen and nitrogen.
- 3. (previously presented) A compound according to claim 1 wherein A is substituted by $-(CH_2)_q \text{aryl or } -(CH_2)_q \text{heteroaryl wherein the aryl or heteroaryl is optionally substituted by one or more substituents independently selected from oxo, C_{1-6}alkyl, halogen, -CN, trifluoromethyl, -OR$^9, -(CH_2)_rCO_2R$^{10}, -NR9R^{10}, -(CH_2)_rCONR9R^{10}, -NHCOR$^9, -SO_2NR9R^{10}, -NHSO_2R$^9 and -S(O)_8R$^9. }$
- 4. (previously presented) A compound according to claim 1 wherein R¹ is methyl.
- 5. (previously presented) A compound according to claim 1 wherein R^2 is -CO-NH-(CH₂)_t- R^{12} .
- 6. (previously presented) A compound according to claim 1 wherein X is hydrogen or fluorine.
- 7. (previously presented) A compound according to claim 1 which is

N-Cyclopropyl-3-[5-fluoro-3-(4-pyridinyl)-1H-indazol-6-yl]-4-methylbenzamide;

N-Cyclopropyl-3-[5-fluoro-3-(1-oxido-4-pyridinyl)-1H-indazol-6-yl]-4-methylbenzamide;

N-Cyclopropyl-3-fluoro-5-[5-fluoro-3-(4-pyridinyl)-1,2-benzisoxazol-6-yl]-4-methylbenzamide;

N-Cyclopropyl-3-fluoro-5-[5-fluoro-3-(1-oxido-4-pyridinyl)-1,2-benzisoxazol-6-yl]-4-methylbenzamide;

 $N-Ethyl-3-\{5-fluoro-3-[6-(methyloxy)-3-pyridinyl]-1H-indazol-6-yl\}-4-methylbenzamide;\\$

3-[3-(6-Chloro-3-pyridinyl)-5-fluoro-1*H*-indazol-6-yl]-*N*-ethyl-4-methylbenzamide;

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or a pharmaceutically acceptable salt thereof.

8. (previously presented) A compound-which is:

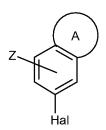
N-cyclopropyl-3-[5-fluoro-3-(4-pyridinyl)-1*H*-indazol-6-yl]-4-methylbenzamide; or N-cyclopropyl-3-fluoro-5-[5-fluoro-3-(4-pyridinyl)-1,2-benzisoxazol-6-yl]-4-methylbenzamide;

or a pharmaceutically acceptable derivative thereof.

9. (previously presented) A pharmaceutical composition comprising at least one compound as claimed in claim 1, or a pharmaceutically acceptable derivative thereof, in association with one or more pharmaceutically acceptable excipients, diluents and/or carriers.

10 to 13. (cancelled)

- 14. (previously presented) A process for preparing a compound of formula (I) as claimed in claim 1, or a pharmaceutically acceptable salt thereof, which comprises
- (a) reacting a compound of formula (II)



(II)

in which A is defined in claim 1 and Hal is halogen, with a compound of formula (IIIA) or (IIIB)

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$$R^1$$
 R^2

(IIIA)

(IIIB)

in which R^1 , R^2 , X and Y are as defined in claim 1, in the presence of a catalyst, or

- (b) final stage modification of one compound of formula (I) as defined in claim 1 to give another compound of formula (I) as defined in claim 1.
- 15. (previously presented) A compound according to claim 3 wherein A is substituted by $-(\text{CH}_2)_q \text{heteroaryl wherein the heteroaryl is optionally substituted by one or more substituents independently selected from 0x0, C_{1-6}alkyl, halogen, -CN, trifluoromethyl, <math display="block"> -\text{OR}^9, -(\text{CH}_2)_r \text{CO}_2 R^{10}, -\text{NR}^9 R^{10}, -(\text{CH}_2)_r \text{CONR}^9 R^{10}, -\text{NHCOR}^9, -\text{SO}_2 \text{NR}^9 R^{10}, -\text{NHSO}_2 R^9 \text{ and } -\text{S(O)}_8 R^9.$
- 16. (previously presented) A compound according to claim 15 wherein R¹ is methyl.
- 17. (previously presented) A compound according to claim 15 wherein R^2 is -CO-NH-(CH₂)_t-R¹².

18. (previously presented) A compound according to claim 15 wherein X is hydrogen or

fluorine.

19 (Currently amended). A compound according to Claim 15 wherein the 5-membered

ring fused to the phenyl ring is an optionally substituted indazole.

20. (previously presented) A compound according to Claim 15 wherein the heteroaryl is a

5- or 6-membered heteroaryl ring containing up to two heteroatoms independently selected

from oxygen and nitrogen.

21. (previously presented) A compound according to Claim 20 wherein the heteroaryl ring

is a pyridyl.

22. (previously presented) A compound according to Claim 21 wherein q is 0.

23. (previously presented) A compound according to Claim 1 wherein Z is a halogen.

24 (previously presented). A compound according to Claim 1 wherein the 5-membered

ring A fused to the phenyl ring is an optionally substituted isoxazolyl, indazole, pyrazolyl or

pyrrolyl.